Abstract

The present invention relates to pyridinone derivatives of formula (I):

$$Z$$
 OH
 OR^1
 OR^1
 OR^1

5

wherein Z represents C_{2-6} alkynyl, aryl or heteroaryl, any of which groups may be optionally substituted, and R^1 represents hydrogen, C_{1-6} alkyl, C_{3-7} heterocycloalkyl(C_{1-6})alkyl, di(C_{1-6})alkylamino(C_{1-6})alkyl,

 C_{2-6} alkylcarbonyloxy(C_{1-6})alkyl or C_{3-7} cycloalkoxycarbonyloxy(C_{1-6})alkyl, and pharmaceutically acceptable salts thereof, useful in the prevention and treatment of hepatitis C virus infections.